

This Page Is Inserted by IFW Operations  
and is not a part of the Official Record

## **BEST AVAILABLE IMAGES**

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

**IMAGES ARE BEST AVAILABLE COPY.**

**As rescanning documents *will not* correct images,  
please do not report the images to the  
Image Problem Mailbox.**

# (12) UK Patent Application (19) GB (11) 2 290 052 (13) A

(43) Date of A Publication 13.12.1995

(21) Application No 9511303.1

(22) Date of Filing 05.06.1995

(30) Priority Data

(31) 08254207 (32) 06.06.1994 (33) US

(71) Applicant(s)

**Kimberly-Clark Corporation**

(Incorporated in USA - Delaware)

**401 North Lake Street, Neenah, Wisconsin 54948,  
United States of America**

(72) Inventor(s)

**Ann Louise McCormack**

**Lance James Garrett Jr**

**Karen Lynn English**

(74) Agent and/or Address for Service

**Frank B Dehn & Co**

**Imperial House, 15-19 Kingsway, LONDON,  
WC2B 6UZ, United Kingdom**

(51) INT CL<sup>6</sup>

**B32B 27/12 // A41D 13/12 31/02, A61F 13/15**

(52) UK CL (Edition N )

**B5N N175 N177 N178 N18X N195 N198 N207 N21Y  
N222 N223 N255 N259 N26X N260 N261 N262 N2702  
N2712 N2732 N2736 N37X N372 N374 N377 N38X  
N383 N389 N39Y N398 N401 N402 N403 N412 N415  
N418 N42X N420 N426 N449 N451 N455 N456 N457  
N46X N466 N491 N501 N502 N518 N537 N577 N578  
N58X N58Y N591 N593 N595 N597 N603 N611 N62X  
N62Y N639 N648 N650 N657 N658 N66Y N661 N662  
N666 N670 N672 N690 N70X N708 N709 N71Y N712  
N734 N738 N764 N766 N767 N77X N778 N78X N784  
U1S S1049 S1122 S1146 S1217 S1218 S1713 S1734  
S1814 S1820 S1859 S2123**

(56) Documents Cited

**GB 2285408 A EP 0309073 A2 WO 93/21013 A1**

(58) Field of Search

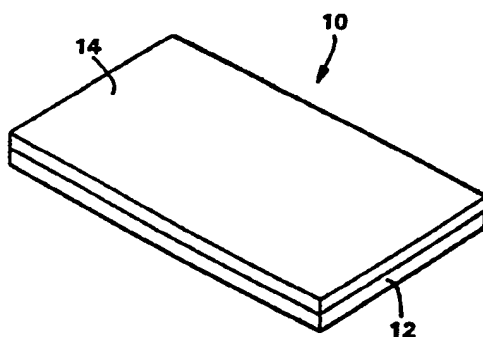
**UK CL (Edition N ) B5N**

**INT CL<sup>6</sup> B32B 27/12**

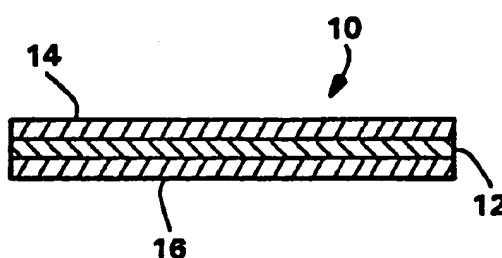
**ONLINE:WPLCLAIMS**

(54) Stretched-thinned film and nonwoven laminate

(57) In a stretch-thinned film and nonwoven laminate (10) the film (12) is polyolefin based and has an effective gauge of 0.55 mils (0.00055 inches) or less and a cross-machine direction to machine direction ratio of Elmendorf tear strength of 3.5 or greater. The resultant laminate has a wide variety of uses which include components of health care related to items such as sterile wrap, surgical drapes and gowns and body side liners, external covers for personal care absorbent products, packaging materials, shower curtains, tents and covers for such items as furniture, computers, automobiles and other vehicles.



**FIG. 1**



**FIG. 2**

**GB 2 290 052 A**

1/2

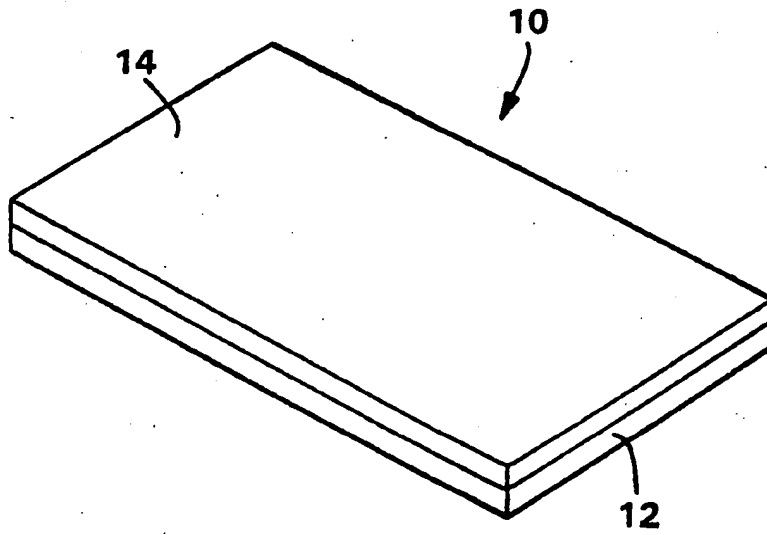


FIG. 1

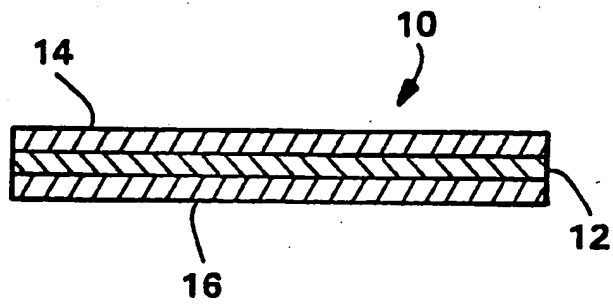


FIG. 2

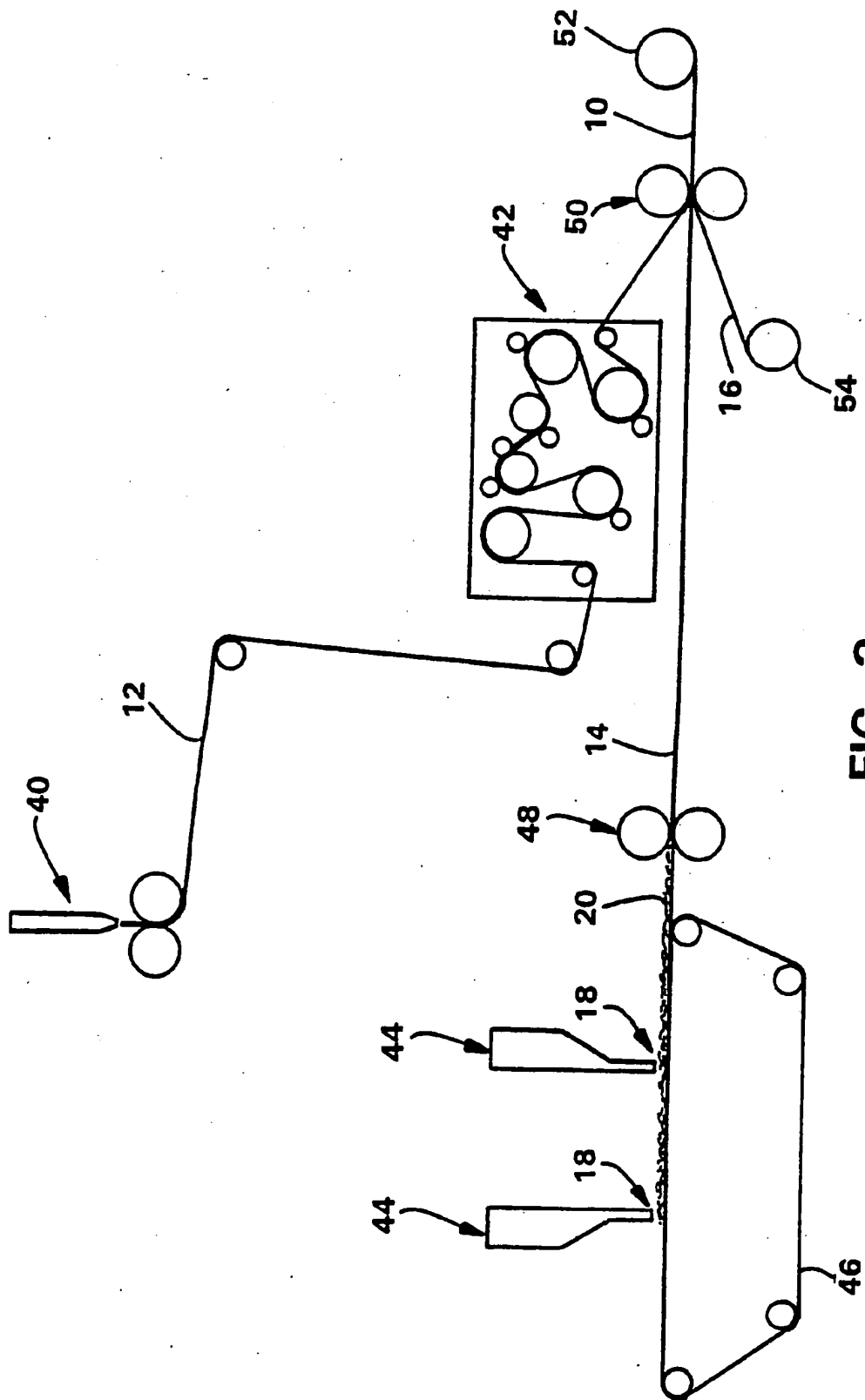


FIG. 3

Stretch-Thinned Film and Nonwoven Laminate

5       The present invention is directed to stretch-  
thinned film and nonwoven laminates. More particularly,  
the present invention is directed to a process for  
making lightweight, low cost, cloth-like and optionally  
breathable film and nonwoven laminates wherein the film  
layer has been thinned to a very low gauge in the  
10   machine direction and subsequently laminated to one or  
more fibrous nonwoven web support layers to create a  
lightweight laminate which has a variety of uses  
including, but not limited to, a component or components  
for personal care absorbent products, articles of  
15   clothing and healthcare related items such as drapes,  
gowns and sterile wrap.

Films have been traditionally used to provide  
barrier properties in single-use items including, but  
not limited to, articles of clothing in general,  
20   protective apparel, healthcare related products  
including surgical drapes, gowns and sterile wrap and  
personal care absorbent products such as diapers,  
training pants, incontinence garments, sanitary napkins,  
bandages and the like. In personal care absorbent  
25   products such as infant diapers and adult incontinence  
products, films are used as the outer covers with the  
purpose of preventing body wastes from contaminating the  
clothing, bedding and other aspects of the surrounding  
environment of use. In the area of protective apparel  
30   including hospital gowns, films are used to prevent  
cross exchange of microorganisms between the wearer and  
the patient. These stand-alone films are usually one to  
two mils in thickness and weigh approximately 0.7 to 1.5  
ounces per square yard. Polyolefin films are most  
35   commonly used in such areas and in their most basic  
forms have a cost of approximately four to six cents per  
square yard based upon a one mil thick film.

While these films can be effective barriers, they are not aesthetically pleasing because their surfaces are smooth and either reel slick or tacky. They are also visually flat and "plasticity" thereby making them less desirable in apparel applications and other uses where they are in contact with human skin. It would be more preferable if these items were more cloth-like from both a tactile and visual standpoint. For example, if infant diapers were more garment-like on their exterior surfaces they could obviate the need for additional clothing. Garment-like adult incontinence products could improve the self image of the incontinent individual. In addition, more garment-like isolation gowns would help the hospital environment feel less foreign and threatening to the patient.

Traditional non-breathable olefin films are available in lighter weights and are usually lower in cost per pound than the microporous films. While weights of 0.7 to 1.5 osy are common, weights as low as 0.4 osy are available. These lightweight films are often difficult to handle and do not have good strength properties.

Laminations of films and nonwovens have been used to create materials which are both impervious and somewhat cloth-like in appearance and texture. The outer covers on premium-priced diapers are but one example. Surgical gowns are another example. Such laminations are more expensive than stand-alone films and their usage has been limited to premium-priced products. Consequently, the need for more cost-effective, cloth-like, impervious barrier laminations has not been fully satisfied.

One reason such existing laminates are more costly than stand-alone films is because more materials are used, thereby resulting in heavier weight products. A second factor is the cost of the lamination process itself. Consequently, the ability to use lightweight

materials and a simple, low-cost lamination process are desirable objectives.

5 A primary purpose of the film in such laminations is to provide barrier properties. In theory, extremely low weights (gauges) of film can meet this requirement. An object of the present invention is to provide extremely low weights and gauges of film in a nonwoven lamination that also can provide the aesthetic and durability properties desired in product usage. There is also a need for such laminates to be breathable so that they have the ability to transmit moisture vapor. Apparel made from laminations of these breathable or microporous films are more comfortable to wear and help keep the wearer's skin healthy by reducing the relative humidity underneath the apparel item. The more traditional, nonbreathable films, however, may be preferred in protective apparel applications where chemical vapor penetration presents a contamination risk to the wearer.

20 An alternative to laminating a film to a nonwoven is to coat the nonwoven with the filming formulation. This process is known as extrusion coating. The extrusion coating process is capable of achieving low weight film coatings. The process relies on the residual heat within the thermally extruded film for adhesion to the nonwoven substrate. One problem with extrusion coating of extremely low weights of film is that the reduced polymer mass carries less heat and makes achieving good adhesion difficult. If adequate adhesion is not achieved, the coating tends to separate from the nonwoven during use so that it can become easily torn or ruptured. The extrusion coated film is not particularly strong since it has not been subjected to an orientation process. Film orientation processes are often used to increase the strength of stand-alone films.

When adequate adhesion of the film and nonwoven is

achieved, the problem of durability during use is largely obviated. However, the structure is somewhat rigid and does not provide the tactile aesthetics that are desired in many applications where a low cost, cloth-like barrier laminate would be appropriate. With extrusion coated constructions a major reason for the stiffness of the overall laminate is the full area adhesion between the two layers. Full area bonding or adhesion provides more durable lamination but results in undesirable stiffness which tends to immobilize the fibers in the nonwoven so that it is not as pleasing to the touch. Consequently, there is a need to balance lamination strength and softness of the overall laminate.

Films may be down-gauged, that is reduced in weight and thickness, by drawing them in either the machine or cross direction. The film is usually heated prior to these drawing processes to make the film more plastic or malleable. This drawing or stretching also orients the molecular structure within the film which increases its strength and durability. The effect of down gauging is desired to reduce cost and the molecular orientation is desired to improve durability. Unfortunately, there are two problems with using stretch-thinned or oriented films far low-cost, cloth-like barriers. The stretch-thinning is an additional process that adds cost to the raw material. Stretching the film in the cross direction is particularly challenging because forces must be applied to the edges of the film to cause it to elongate. Tenter frames are commonly used. In contrast, stretch-thinning the film in the machine direction is relatively easy. It is only necessary to increase the draw, or speed ratio, between two rollers while the film is in the heated or plastic state. Machine-directionally stretched films are desired to achieve the lowest costs in down-gauged films because of process simplicity and the process speeds that can be



achieved in comparison to cross-directionally stretch-thinned films.

5       There is a durability problem, however, with uni-directionally-stretch-thinned films, be it machine direction or cross-direction. Uni-directional stretching can achieve the down gauging objectives, but the molecular orientation occurs only in the stretched direction. This results in films that are easily torn or split along that dimension. For example, a machine-directionally oriented film has a propensity to split or  
10       tear along the machine direction. Also, the tensile characteristics of the film are dramatically increased in the machine direction, but the tensile strength in the cross-direction is significantly inferior to that of  
15       the machine direction.

      These durability problems with uni-directionally stretched or oriented films are well known. Two approaches are commonly used to obviate the product durability problems resulting from these highly  
20       isotropic strength characteristics. The first is to stretch-orient the film in both the machine and cross direction. Films that have been biaxially stretched have more balanced strength properties. The second approach is to combine into a laminate one layer of  
25       machine directionally oriented film with one layer of cross-directionally oriented film. Neither approach is consistent with the cost objectives of the present invention because both entail the use of the cross directional stretching process. Further, the second  
30       approach doubles the minimum weight that can be achieved in the final laminate and consequently the cost of the material. There is therefore a need for a machine-directionally-stretched, lightweight film and nonwoven laminated using low-cost materials and processes that  
35       provides a laminate with both the cloth-like aesthetics and the in-use durability that are desired.

      The present invention is directed to a stretch-

thinned film and nonwoven laminate and a process for making such a laminate. The process involves sufficiently stretching a polyolefin-based film to cause the film to have an effective gauge of 0.55 mils or less and a cross-machine direction to machine direction ratio of Elmendorf tear strengths of 3.5 or greater. Next a first fibrous nonwoven web support layer is bonded to the film to form the laminate. Desirably, the bonding of the laminate is in the form of discrete bond points with a maximum total bond area of about 60 percent. Optionally, a second fibrous nonwoven web support layer may be bonded to a surface of the film layer opposite that which is bonded to the first fibrous nonwoven web support layer.

The resultant two layer laminate has a ratio of cross-machine direction to machine direction Elmendorf tear strengths of less than 3.5, preferably with the minimum machine direction Elmendorf tear strength being 100 grams or greater and the peak load cup crush value being less than 150 grams. With the three layer laminate the softness will not be as good but in any event the peak load cup crush value should preferably be 300 grams or less.

Embodiments of the invention will now be described by way of example only and with reference to the accompanying drawings, in which:

Figure 1 is a perspective view of a stretch-thinned film and nonwoven laminate according to the present invention.

Figure 2 is a cross-sectional side view of another stretch-thinned film and nonwoven laminate according to the present invention.

Figure 3 is a schematic side view of a process for forming a stretch-thinned film and nonwoven laminate according to the present invention.

Referring to Fig. 1, there is shown a stretch-

thinned film and nonwoven laminate 10 according to the present invention including a first film layer 12 and a first fibrous nonwoven web support layer 14. The film layer 12 can be made from either cast or blown film equipment and can be embossed if so desired. The film layer 12 can be made from any thermoplastic film which can be stretched in one direction such that the film gauge or thickness can be reduced from an initial gauge to an effective final gauge of 0.55 mils or less. Generally, this stretching will occur in the machine direction. In addition, stretching may take place in the cross-machine direction provided, however, that the ratio of the cross-machine direction (CD) Elmendorf tear strength to the machine direction (MD) Elmendorf tear strength does not fall below about 3.5.

In some applications it may be desirable to make the overall laminate breathable. While most nonwovens are breathable, the same is not true with respect to films. Some films are made breathable by adding filler particles such as calcium carbonate to the film during the film-forming process. Once the particle-filled film has been formed, it is then either stretched or crushed to create pathways through the film, thereby making the film breathable. This too can add an additional step to the overall process and increase the price of the resultant material. Other films, such as a film sold under the trade designation FX-1801 Scotchban Protector Film from Minnesota Mining and Mineral Company of St. Paul, Minnesota, are available which are microporous and thus breathable without the use of fillers. Generally to qualify as being "breathable" for the present application, the resultant film/nonwoven laminate should have a water vapor transmission rate of 200 grams per square meter per 24 hours or greater as defined by the test method described below.

Suitable polymers for forming the film layer 12 are

any polymers or polymer blends which can be formed into films, stretch-thinned to an effective gauge or 0.55 mils or less and then be bonded to a fibrous nonwoven support layer 14 without causing pinholes in the film layer which would permit the passage of liquids such as water, urine, blood and other body fluids. Polyolefin-based films have been found to work particularly well with respect to the present invention. For purposes of the present invention, a film is considered to be "polyolefin-based" if the polymer portion of the film, exclusive of any filler materials, has at least 50 weight percent polyolefin based upon the total weight of polymer in the film.

In addition to polymer, the film layer 12 may also include a filler to make the film breathable, more opaque or to impart some other property. As used herein a "filler" is meant to include particulates and other forms of materials which can be added to the film polymer extrusion blend and which will not chemically interfere with or adversely affect the extruded film but which are able to be uniformly dispersed throughout the film. Generally the fillers will be in particulate form and usually will have somewhat of a spherical shape with average particle sizes in the range of about 0.1 to about 7 microns. In addition, the film will usually contain at least 30 percent filler based upon the total weight of the film layer. Both organic and inorganic fillers are contemplated to be within the scope of the present invention provided that they do not interfere with the film formation process, the breathability of the resultant film or its ability to bond to the fibrous polyolefin nonwoven web or comfort layer. Examples of fillers include calcium carbonate ( $\text{CaCO}_3$ ), various kinds of clay, silica ( $\text{SiO}_2$ ), alumina, barium sulfate, sodium carbonate, talc, magnesium sulfate, titanium dioxide, zeolites, aluminum sulfate, cellulose-type powders,

diatomaceous earth, magnesium sulfate, magnesium carbonate, barium carbonate, kaolin, mica, carbon, calcium oxide, magnesium oxide, aluminum hydroxide, pulp powder, wood powder, cellulose derivative, polymer particles, chitin and chitin derivatives.

An important aspect of the present invention is the ability to form a very thin film with sufficient strength in the machine direction and then couple it with a fibrous nonwoven support layer to yield a laminate which is low cost due to maximum reduction in materials and at the same time, sufficiently soft to the touch and strong so as to be suitable for uses such as the outer cover of a personal care absorbent article. Consequently, applicants have found that the film layer 12 must have an "effective" film thickness or gauge of 0.55 mils or less. The effective gauge is used to take into consideration the voids or air space in breathable film layers. For normal non-filled/non-breathable films the actual gauge and the effective gauge of the film will most likely be the same. However, for filled films that have been stretched, the thickness of the film will also include the air spaces. In order to disregard this added volume, the effective thickness is calculated according to the test method set forth below.

Once a film has been stretch-thinned to an effective gauge of 0.55 mils or less, it will tend to be "splitty." By splitty it is meant that the film will tend to split or tear along lines parallel to the machine direction (the direction of stretching), thereby making it weak from a tensile standpoint in the cross-machine direction. As a result, the film must be reinforced by some type of support layer 14. In addition, the overall laminate must be soft to the touch, at least on one side. Consequently, the film layer 12 is bonded to a layer of fibrous nonwoven web material generally having a fairly low basis weight such as two ounces per square yard (osy) and even one osy and

less.

The support layer 14 can be any material which will provide a soft feel and a laminate with a ratio of cross-machine direction to machine-direction Elmendorf  
5 tear strengths of less than 3.5. Fibrous nonwoven webs have been found to work particularly well due to their low cost, good strength, integrity and soft feel. Spunbond webs which have long essentially continuous fibers, once bonded and properly engineered, work well.  
10 The formation of spunbond webs is well-known. See for example patents such as Appel et al., U.S. Patent 4,340,562; Dorschner et al., U.S. Patent 3,692,618; Kinney, U.S. Patent Numbers 3,338,992 and 3,341,394; Levy, U.S. Patent Number 3,276,944; Peterson, U.S.  
15 Patent 3,502,538; Hartman, U.S. Patent 3,502,763, Matsuki et al., U.S. Patent 3,802,817 and Dobo et al., U.S. Patent Number 3,542,615 all of which are incorporated herein by reference in their entirety.

Suitable polymers for forming a fibrous nonwoven  
20 web support layer include, but are not limited to, homopolymers, copolymers and blends of polyolefins and polyesters. In addition, the fibers may be single component fibers or multicomponent fibers such as bicomponent fibers and biconstituent fibers.  
25 Biconstituent fibers are extruded from a homogeneous mixture of two different polymers. Such fibers combine the characteristics of the two polymers into a single fiber. Bicomponent or composite fibers are composed of two or more polymer types in distinct areas of the fiber  
30 such as in a side-by-side or sheath-core configuration.

Fiber size and basis weight can be varied, bearing in mind the requirements of softness and strength. The degree of bonding of the web will also affect these  
35 properties. Polymer selection will also affect softness. Generally, fiber sizes will be about 6 denier or less and basis weights will be about two ounces per

square yard or less. In heavy duty applications, the basis weights may be higher. Bonding of the fibers in the support layer 14 can be through the use of heat and/or pressure as with ultrasonic bonding equipment and heated bonding rolls. Point bonding has been found to create a softer nonwoven web than overall bonding with total bond area of 20 percent or less being desirable. Other types of bonding can include adhesive bonding such as with latexes, powdered or dry adhesives and solvent-based adhesives.

Bonding or lamination of the film layer 12 to the support layer 14 can be by any suitable means which will yield a laminate with sufficient MD strength as measured by its Elmendorf tear strength of 100 grams or greater and sufficient softness as measured by its cup crush value of less than 150 grams. In addition, where liquid barrier properties are important, the laminating process should not result in a laminate that will leak. In view of the foregoing requirements, possible laminating techniques or methods include, but are not limited to, ultrasonic bonding, adhesive bonding and thermomechanical bonding such as with heated and patterned laminating rolls. Lamination may take place across the entire interface between the two layers, however, extensive bonding tends to reduce the overall softness of the laminate 10. As a result, relative to a single side surface of the laminate, it has been found to be more desirable to limit the total bond area relative to the surface area being measured to a maximum of 60 percent. Generally this can be accomplished by using a bond pattern having discrete bond points or by applying discrete sites of adhesion.

As described thus far, the laminate 10 has included a film layer 12 and a single support layer 14 laminated to one side of the film layer 12. If desired, it is also possible to laminate a second support layer 16 to a side of the film layer 12 opposite the first support

layer 14. While not being necessary, it is most likely that the second support layer 16 will also be a fibrous nonwoven web. As a result, the options and features described previously with respect to the first support layer 14 can also be employed with respect to the second support layer 16. In addition, the same types of laminating techniques described above may be used in the lamination of the second support layer 1,6 to the film layer 12.

Creating a three-layer laminate in lieu of a two-layer laminate can provide certain advantages, especially with respect to certain end-use applications. Providing a second support layer 16 can improve the overall strength of the laminate 10, especially in the MD direction. It should be appreciated, however, that with increased overall basis weight for the laminate, cup crush values may increase though it is still possible to achieve peak load cup crush values of 200 grams or less.

Referring to Figure 3, there is shown in schematic form, a process for forming a stretch-thinned film and nonwoven laminate according to the present invention. The film layer 12 is formed using any type of conventional film forming equipment 40 such as cast or blown equipment. As mentioned earlier, the film 12 so formed, if so desired, can contain a filler or other additive to impart particular properties to the film layer 12. After the film layer 12 has been formed, it is sent through a film stretching apparatus 42 to stretch and thin the film to an effective gauge of 0.55 mils or less. One type of apparatus 42 suitable for such stretch-thinning is a Machine Direction Orienter (NDO) Unit, Model No. 7200 from the Marshall and Williams Company of Providence, Rhode Island.

While the film layer 12 is being formed and thinned, the fibrous nonwoven web support layer 14 is also being formed. Referring to Figure 3, a conventional



fibrous nonwoven web forming apparatus 44, such as a spunbond machine, is used to form a support layer 14. The long, essentially continuous fibers 18 are deposited onto a forming wire 46 as an unbonded matt 20 and the  
5 unbonded matt is then sent through a pair of bonding rolls 48 to bond the fibers together and increase the tear strength of the resultant web 14. One or both of the rolls are often heated to aid in bonding. Typically, one of the rolls 48 is also patterned so as  
10 to impart a discrete bond pattern with a prescribed bond surface area to the web 14. The other roll is usually a smooth anvil roll but this roll may also be patterned if so desired.

Once the film layer 12 has been sufficiently  
15 thinned and oriented and the support layer 14 has been formed, the two layers are brought together and laminated to one another using a pair of laminating rolls or other means 50. As with the bonding rolls 48, the laminating rolls 50 may be heated. Also, at least  
20 one of the rolls is patterned to create a discrete bond pattern with a prescribed bond surface area for the laminate 10. Generally, the maximum bond point surface area for a given area of surface on one side of the laminate will not exceed about 60 percent of the total  
25 surface area.

After the laminate has been formed, it is wound into a roll 52 for subsequent processing. Alternatively, the laminate 10 may continue in-line for further conversion.

30 It should be noted that the process shown in Figure 3 may be altered in a number of ways without departing from the spirit and scope of the present invention. For example, a different apparatus 42 can be used for stretch-thinning the film layer 12. Different  
35 film/nonwoven web forming equipment 44 such as meltblown and bonded carded web equipment may be used in place of the spunbond equipment. In addition, other means for

bonding and laminating the support layer 14 and laminate 10 may be used provided the resultant laminate 10 has the required properties described herein. Lastly, the formation processes for making the film and support layers may be done at a remote site and rolls of the two materials may be unwound into the process.

Referring again to Figure 3, the process shown also may be used to create a three layer laminate 10 such as is shown in Figure 2 of the drawings. The only modification to the previously described process is to feed a supply 54 of a second fibrous nonwoven web support layer material 16 into the laminating means 50 on a side of the film layer 12 opposite that of the first fibrous nonwoven web support layer 14. As shown in Figure 3, the supply of material for layer 16 is in the form of a preformed roll 54 of material. As with the other layers 12 and 14, layer 16 may be formed directly in-line or it may be preformed and then fed into the process from a supply roll. In either event, the second support layer 16 is fed into the laminating means 50 and is laminated to the film layer 12 in the same fashion as the first support layer 14.

#### TEST METHODS

The properties of the present invention were determined using a series of test procedures. These properties included MD and CD Elmendorf tear strengths, cup crush which is related to softness, effective gauge and water vapor transmission rate. The test procedures and/or a reference to the published standards are set forth below. Three replications were performed for each of the test values given in the examples.

#### Elmendorf Tear Strength

The Elmendorf tear strength is a measure of the

average force required to propagate a single-nip, tongue-type tear starting from a cut in a fabric by means of a falling-pendulum (Elmendorf) apparatus. It can be measured in both the machine direction (MD) and cross-machine direction (CD) for both single layer materials and laminates. Samples were cut having dimensions of 102 millimeters by 63 millimeters. For cross-machine direction measurements, samples were cut with the long axis of the sample being parallel to the machine direction of the sample and thus the direction of orientation. For machine direction measurements samples were cut with the long axis of the sample being perpendicular to the machine direction of the sample and thus the direction of orientation. In all cases, the initial cut in the samples was in the longer side of the specimens. As shown by the data below, the CD Elmendorf tear strengths were greater than the MD values. The test is performed in accordance with ASTM Standard Test Method D1424-83 except that the 6 millimeter deviation rule in section 11.5 of the test procedure was disregarded.

#### Cup Crush

The cup crush of an individual layer or laminate was measured according to the following procedure and is shown in units of grams. The cup crush is an indication of the softness of a material and the lower the cup crush value, the softer the material. This procedure was conducted in a controlled environment wherein the temperature was about 73°F and the relative humidity was about 50 percent. Samples were tested using a Material Test Instrument and Crush Test Stand available from Kimberly-Clark Corporation Quality Assurance Department in Neenah, Wisconsin. The Material Test Instrument and Crush Test Stand included a model 11 foot, a model 21, forming cylinder, a model 31, steel ring, a model 41,

forming cup, a calibration set and an Epson FX-86e printer with cable.

5       The steel ring was placed over the forming cylinder and a 9 x 9 inch sample was centered over the forming cylinder. The forming cylinder was inserted into the forming cup until the sample was pinched between the forming cylinder and the steel ring all the way around the steel ring. The forming cup was placed on top of the cylinder plate of the load cell and firmly seated  
10       over the ridge of the cylinder plate. The foot was mechanically lowered into the forming cup, crushing the sample while the Materials Test Instrument measured the peak load needed to crush the sample.

15       Water Vapor Transmission Rate

          The water vapor transmission rate (WVTR) for the sample materials was calculated in accordance with ASTM Standard E96-80. Circular samples measuring three  
20       inches in diameter were cut from each of the test materials and a control which was a piece of CELGUARD® 2500 film from Hoechst Celanese Corporation of Sommerville, New Jersey. CELGUARD® 2500 film is a microporous polypropylene film. Three samples were  
25       prepared for each material. The test dish was a number 60-1, Vapometer pan distributed by Thwing-Albert Instrument Company of Philadelphia, Pennsylvania. One hundred millileters of water was poured into each Vapometer pan and individual samples of the test  
30       materials and control material were placed across the open tops of the individual pans. Screw-on flanges were tightened to form a seal along the edges of each pan, leaving the associated test material or control material exposed to the ambient atmosphere over a 6.5 centimeter  
35       diameter circle having an exposed area of approximately 33.17 square centimeters. The pans were placed in a forced air oven at 100°F (32°C) for 1, hour to

equilibrate. The oven was a constant temperature oven with external air circulating through it to prevent water vapor accumulation inside. A suitable forced air oven is, for example, a Blue M Power-O-Matic 60 oven distributed by Blue M Electric Company of Blue Island, Illinois. Upon completion of the equilibration, the pans were removed from the oven, weighed and immediately returned to the oven. After 24 hours, the pans were removed from the oven and weighed again. The preliminary test water vapor transmission rate values were calculated as follows:

$$\text{Test WVTR} = (\text{grams weight loss over 24 hours}) \\ \times 315.5 \text{ g/m}^2 / 24 \text{ hrs}$$

The relative humidity within the oven was not specifically controlled.

Under predetermined set conditions of 100°F (32°C) and ambient relative humidity, the WVTR for the CELGUARD® 2500 control has been determined to be 5000 grams per square meter for 24 hours. Accordingly the control sample was run with each test and the preliminary test values were corrected to set conditions using the following equation:

$$\text{WVTR} = (\text{Test WVTR/control WVTR}) \times 5000 \text{ g/m}^2/24 \text{ hrs.}) \\ (\text{g/m}^2/ 24 \text{ hrs})$$

#### Effective Gauge

The effective gauge of a material was calculated by dividing the basis weight of the film layer by the density of the polymer(s) and fillers forming the film.

The effective gauge of a layer of film was calculated by multiplying 0.001334 (a metric to English conversion factor) times the weight per unit area of the film sample in ounces per square yard and dividing the

result by the density of the polymer formulation in grams per cubic centimeter to yield the effective gauge in inches.

5     **EXAMPLES**

**Example I**

      In Example I a two layer, breathable stretch-  
10     thinned film and nonwoven laminate was made according to  
      the present invention. The film layer contained, on a  
      total weight percent basis, 65 percent English China  
      Supercoat calcium carbonate ( $\text{CaCO}_3$ ) with a 1, micron  
      average particle size and a 7 micron top cut. The  
15     calcium carbonate was obtained from ECCA Calcium  
      Products, Inc. in Sylacauga, Alabama, a division of ECC  
      International. The calcium carbonate was blended with  
      20 percent by weight of linear low density polyethylene  
      made from a blend of Dowlex® 2517 linear low density  
20     polyethylene and Dowlex® 2532 linear low density  
      polyethylene blended in a weight ratio of 1:4 such that  
      the melt index of the blend was 10 M.I. (/10 minutes at  
      190°F). The Dowlex® polymers are available from Dow  
      chemical U.S.A., Midland, Michigan. The remaining 15  
25     percent by weight of the formulation comprised Himont  
      KS05IP polypropylene-based polymer from Himont, USA of  
      Wilmington, Delaware. The KS05IP polymer is an olefinic  
      thermoplastic elastomer or TPO multistep reactor product  
      wherein an amorphous ethylene propylene random copolymer  
30     is molecularly dispersed in a predominately  
      semicrystalline high polypropylene monomer/low ethylene  
      monomer continuous matrix. The film had an initial  
      gauge of 1.5 mil and was stretch-thinned to an effective  
      gauge of 0.42 mils using a machine direction orienter  
35     (MDO) unit of the type shown in Figure 3. The film was  
      heated to a temperature of 170°F and the film was run  
      through the MDO unit at a line speed of 500 feet per

minute to stretch the film approximately four times its original length.

5       The first fibrous nonwoven web support layer was a 0.5 ounce per square yard spunbond web made from approximately 2.0 to 2.5 denier polypropylene fibers. The polymer used to make the spunbond web was Exxon 3445 polypropylene from the Exxon Chemical Company of Houston, Texas. The spunbond web was prebonded using discrete bond points with a total bond area of 17 percent per unit area of web.

10       Lamination of the two layers was effected using a patterned laminating roll with a baby objects pattern at a temperature of 230°F and a smooth anvil roll at a temperature of 160°F. The film layer was positioned against the anvil roll and the pressure was set at 20 pounds per square inch gauge (psig). The resultant laminate had total bond area of 14 percent per unit area and a basis weight of 1.15 osy (38.3 grams per square meter).

20       Testing of the laminate indicated that the stretch-thinned film layer by itself had a peak load cup crush value of 75 grams, a CD and MD Elmendorf tear strength of 144 and 32 grams, respectively. The ratio of the CD to MD Elmendorf tear strength was 4.5. The laminate had a peak load cup crush of 76 grams. Elmendorf tear strengths for the laminate in the CD and MD directions were 496 and 224 grams respectively and the CD/MD ratio was 2.2. Lastly, the laminate had a water vapor transmission rate of 3700 grams per square meter per 24 hours.

#### Example II

35       In Example II a two-layer non-breathable, stretch-thinned film and nonwoven laminate was made according to the present invention. The film layer comprised on a weight percent basis, based upon the total weight of the

film, 1.9 percent titanium dioxide ( $\text{TiO}_2$ ) concentrate, 32 percent polypropylene, 5 percent low density polyethylene and 32 percent of a polypropylene-based polymer designated Himont K5080 Catalloy polymer from Himont, USA of Wilmington, Delaware. The titanium dioxide was Ampacet 110310 from Ampacet Corporation of Tarrytown, New York. The polypropylene was Exxon 3445 polypropylene from Exxon Chemical Company of Houston, Texas and the low density polyethylene was Quantum NA334 low density polyethylene from Quantum Chemical Corporation of New York, New York. The film had an initial gauge of 0.6 mils and was stretch-thinned to an effective gauge of 0.41 mils. During the stretch-thinning process the film was heated to a temperature of 190°F and the film was stretched to approximately two times its original length at a line speed of 300 feet per minute. The film layer by itself had a peak load cup crush value of 36 grams and MD and CD Elmendorf tear strengths of 16 and 320 grams, respectively. The fibrous nonwoven web support layer was the same as that used in Example I. Lamination conditions and lamination bond area were the same as in Example I except for the laminating pressure which was 25 psig.

The resultant laminate had a basis weight of 0.8 osy (26.8 grams per square meter), a peak load cup crush value of 104 grams, a CD Elmendorf tear strength of 352 grams and a MD Elmendorf tear strength of 160 grams. The ratio of the CD to MD Elmendorf tear strengths was 2.2.

### Example III

In Example III, a three layer, stretch-thinned film and nonwoven laminate was made according to the present invention. The first and second fibrous nonwoven web support layers were the same 0.5 osy spunbond web material used in Examples I and II and the film layer



was made from the same film material with the same 0.42 mil effective gauge as was used in Example I.

Lamination of the three layers, with the film in the middle, took place using the same patterned and anvil rolls as were used in Examples I and II. As a result, total lamination band area per unit area of either support layer was 14 percent. Lamination took place at a line speed of 100 feet per minute with both the patterned and anvil rolls heated to a temperature of 235°F and with the lamination pressure between the rolls set at 30 psig.

The resultant laminate had a basis weight of 1.6 osy (53.6 grams per square meter), a water vapor transmission rate of 3700 grams per square meter per 24 hours and a peak load cup crush value of 140 grams. CD and ND Elmendorf tear strengths for the laminate were 640 and 224 grams, respectively, and the ratio of CD to MD Elmendorf tear strengths was 2.9.

The stretch-thinned film and nonwoven laminate of the present invention has a wide variety of uses and applications. Non-restrictive examples of such uses and applications include body side liners and outer covers for personal care absorbent products. Typically such products include a body side liner which faces the user and an outer cover which forms the external surface of the product. Disposed between these two materials there is usually an absorbent core for absorbing body exudates such as blood, urine, feces and other body fluids. The material of the present invention may be used to form all or a portion of either or both of the body side liner and the outer cover.

Articles of clothing also sometimes employ films and/or nonwovens in their construction. As a result, the material of the present invention may be used to form all or a portion of such articles.

In the health care arena many surgical drapes, gowns and sterile wraps are made from films and/or

nonwovens. Consequently, the material of the present invention may be used to form all or a portion of any one of these products as well.

5 Other areas of possible applications for the material of the present invention include, but are not limited to, packaging materials, shower curtains, tent material and covers for such items as furniture, computers, automobiles and other vehicles.

10 Having thus described the invention in detail, it should be appreciated that various modifications and changes can be made to the present invention without departing from the scope of the following claims.

Claims

1. A process for forming a stretch-thinned film and nonwoven laminate comprising:

- 5       a) sufficiently stretching a polyolefin-based film to cause said film to have an effective gauge of 0.55 mils or less and a cross-machine direction to machine direction ratio of Elmendorf tear strength of 3.5 or greater, and
- 10       b) bonding a first fibrous nonwoven web support layer to said film to form a laminate.

2. The process of claim 1 wherein said bonding creates a plurality of discrete bond points between said  
15 film and said first fibrous nonwoven web support layer with a maximum total bond area of about 60 percent.

3. The process of claim 1 or 2 which further includes bonding a second fibrous nonwoven web support  
20 layer to a surface of said film opposite said first fibrous nonwoven web support layer.

4. A stretched-thinned film and nonwoven laminate comprising:

- 25       a polyolefin-based film layer having an effective gauge of 0.55 mils or less, a ratio of cross-machine direction to machine direction Elmendorf tear strengths of 3.5 or greater, and
- 30       a first fibrous nonwoven web support layer bonded to said film layer.

5. The stretch-thinned film and nonwoven laminate of claim wherein said film layer is bonded to said first  
35 fibrous nonwoven web support layer at a plurality of discrete bond points with a maximum total bond area of about 60 percent.

6. The stretched-thinned film and nonwoven laminate of claim 4 or 5 wher in a second fibrous nonwoven web support layer is bonded to a surface of said film layer opposite said first fibrous nonwoven web support layer.

7. The stretched-thinned film and nonwoven laminate of claim 4 or 5 wherein said laminate has a minimum machine direction Elmendorf tear strength of 100 grams or greater and a peak load cup crush value of less than 150 grams.

8. The stretched-thinned film and nonwoven laminate of claim 6 wherein said laminate has a peak load cup crush value of 300 grams or less.

9. The stretch-thinned film and nonwoven laminate of any of claims 4 to 8 wherein said laminate has a water vapor transmission rate of 300 grams/meter<sup>2</sup>/24 hours or greater.

10. The stretch-thinned film and nonwoven laminate of any of claims 4 to 9 wherein said film contains at least 30 percent filler based upon the total weight of said film layer.

11. The stretch-thinned film and nonwoven laminate of any of claims 4 to 10 wherein said laminate has a ratio of cross-machine direction to machine direction Elmendorf strengths of less than 3.5.

12. A personal care absorbent product comprising a body side liner, an outer cover and an absorbent core disposed between said body side liner and said outer cover, at least one of said body side liner and said outer cover comprising the stretch-thinned film and nonwoven laminate of any of claims 4 to 11.

13. An article of clothing wherein at least a portion of said article comprises the stretch-thinned film and nonwoven laminate of any of claims 4 to 11.

5           14. A surgical drape wherein at least a portion of said drape comprises the stretch-thinned film of nonwoven laminate of any of claims 4 to 11.

10           15. A sterile wrap wherein at least a portion of said wrap comprises the stretch-thinned film and nonwoven laminate of any of claims 4 to 11.

15           16. A process for forming a stretch-thinned film and nonwoven laminate substantially as hereinbefore described with reference to Figs. 1 and 3 or Figs. 2 and 3 of the accompanying drawings.

20           17. A stretch-thinned film and nonwoven laminate substantially as hereinbefore described with reference to Figs. 1 and 3 or Figs. 2 and 3 of the accompanying drawings.

**Patents Act 1977**  
**Examiner's report to the Comptroller under Section 17**  
**(T Search report)**

26

Application number  
GB 9511303.1

**Relevant Technical Fields**

- (i) UK Cl (Ed.N) B5N  
(ii) Int Cl (Ed.6) B32B 27/12

Search Examiner  
R J MIRAMS

Date of completion of Search  
19 SEPTEMBER 1995

**Databases (see below)**

- (i) UK Patent Office collections of GB, EP, WO and US patent specifications.

Documents considered relevant following a search in respect of Claims :-  
1 TO 17

- (ii) ONLINE: WPI, CLAIMS

**Categories of documents**

- X: Document indicating lack of novelty or of inventive step. P: Document published on or after the declared priority date but before the filing date of the present application.  
Y: Document indicating lack of inventive step if combined with one or more other documents of the same category. E: Patent document published on or after, but with priority date earlier than, the filing date of the present application.  
A: Document indicating technological background and/or state of the art. &: Member of the same patent family; corresponding document.

Category	Identity of document and relevant passages	Relevant to claim(s)
X	GB 2285408 A (KIMBERLY-CLARK) whole document	at least 1, 4, 5, 9, 10 and 12 to 15
A	EP 0309073 A2 (EXXON)	
A	WO 93/21013 A1 (W L GORE)	

Databases: The UK Patent Office database comprises classified collections of GB, EP, WO and US patent specifications as outlined periodically in the Official Journal (Patents). The on-line databases considered for search are also listed periodically in the Official Journal (Patents).